## **Amendments to the Specification:**

Please replace the 1st paragraph of page 1 with the following rewritten paragraphs:

# Reference to Related Applications

This is a divisional of <del>copending applications Serial number</del> <u>U.S.S.N.</u> 08/400,796 filed on March 7, 1995, now <u>U.S. Pat. No. 5,874,531</u>.

#### Amendments to the Specification:

This listing of claims will replace all prior versions, and listings, of claims in the application:

#### 1-2. (Cancelled)

3. (Currently amended) A pharmaceutical preparation for tolerizing, comprising a pharmaceutically acceptable carrier and

an amount of an isolated human polypeptide effective for tolerizing an individual to an autoantigen, said human polypeptide consisting of a sequence motif for an HLA-DR protein containing the core MHC binding residues, wherein said HLA-DR protein is selected from the group consisting of HLA-DR2 and HLA-DR4 consisting essentially of an amino acid sequence corresponding to the core MHC binding residues of a sequence motif for an HLA-DR protein;

wherein said sequence motif for said HLA-DR protein is based upon the structure of the HLA-DR peptide binding site;

wherein said HLA-DR protein is associated with a human autoimmune disease <u>selected</u> from <u>Pemphigus Vulgaris (PV)</u> and <u>Multiple Sclerosis (MS)</u>;

wherein said polypeptide binds to said HLA-DR protein;

wherein said polypeptide bound to said HLA-DR protein activates autoreactive T cells from a subject having said autoimmune disease; and

wherein said protein is a non-collagen and non-myelin basic protein polypeptide.

- 4. (**Original**) The pharmaceutical preparation of claim 3 wherein said HLA-DR protein is an HLA-DR4 protein and said autoimmune disease is pemphigus vulgaris.
- 5. (Original) The pharmaceutical preparation of claim 4 wherein said motif is PV motif #1.
- 6. (Currently amended) The pharmaceutical preparation of claim 4, wherein said amino acid sequence polypeptide consists essentially of an amino acid sequence selected from the group consisting of SEQ ID NO:: 1, SEQ ID NO:: 2, SEQ ID NO:: 3, SEQ ID NO:: 4, SEQ ID NO:: 5, SEQ ID NO:: 6, and or SEQ ID NO:: 7.

### 7-10. (Cancelled)

11. (Original) A method of tolerizing an individual to an autoantigen of pemphigus vulgaris comprising administering an effective amount of the pharmaceutical preparation of any one of claims 4-6 to a subject in need of such treatment.

#### 12. (Cancelled)

13. (Currently amended) A pharmaceutical preparation for vaccinating an individual at risk of an autoimmune disease, comprising a pharmaceutically acceptable carrier and an amount of an immunogenic preparation antigen from a human pathogen, and effective to immunize against the a human pathogen, wherein said antigen does not comprise a sequence motif for an HLA-DR protein containing the core MHC binding residues, wherein said HLA-DR protein is selected from the group consisting of HLA-DR2 and HLA-DR4 that in its native form includes a polypeptide having an amino acid sequence corresponding to the core MHC binding residues of a sequence motif for an HLA-DR protein;

wherein said sequence motif for said HLA-DR protein is based upon the structure of the HLA-DR peptide binding site;

wherein said HLA-DR protein is associated with a human autoimmune disease <u>selected</u> from Pemphigus Vulgaris (PV) and Multiple Sclerosis (MS);

wherein said polypeptide binds to said HLA-DR protein; and

wherein said polypeptide bound to said HLA-DR protein activates autoreactive T cells from a subject having said autoimmune disease; and

wherein said preparation is free of a polypeptide corresponding to said sequence.

- 14. (Original) The pharmaceutical preparation of claim 13 wherein said HLA-DR protein is an HLA-DR4 protein and said autoimmune disease is pemphigus vulgaris.
- 15. (Original) The pharmaceutical preparation of claim 14 wherein said motif is PV motif #1.

16. (Currently Amended) The pharmaceutical preparation of claim 14 wherein said amino acid sequence polypeptide consists essentially of an amino acid sequence selected from the group consisting of SEQ ID NO-: 1, SEQ ID NO-: 2, SEQ ID NO-: 3, SEQ ID NO-: 4, SEQ ID NO-: 5, SEQ ID NO-: 6, and or SEQ ID NO-: 7.

17-29. (Cancelled)